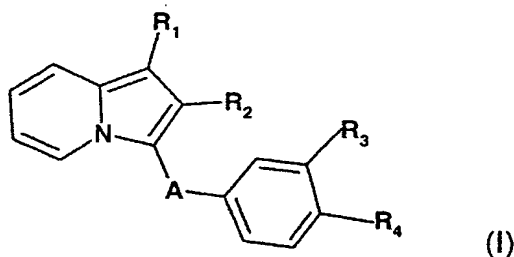


Novel 1,2,3-substituted indolizine derivatives,  
inhibitors of FGFs, method for preparing them and  
pharmaceutical compositions containing them

ABSTRACT

The present invention relates to derivatives  
of formula I:



in which

- $R_1$  represents -OH,  $(C_1-C_5)$ alkoxy, carboxyl,  $(C_2-C_6)$ -alkoxycarbonyl,  $-NR_5R_6$ ,  $-NH-SO_2-Alk$ ,  $-NH-SO_2-Ph$ ,  $-NH-CO-Ph$ ,  $-N(Alk)-CO-Ph$ ,  $-NH-CO-NH-Ph$ ,  $-NH-CO-Alk$ ,  $-NH-CO_2-Alk$ ,  $-O-(CH_2)_n-cAlk$ ,  $-O-Alk-COOR_7$ ,  $-O-Alk-O-R_8$ ,  $-O-Alk-OH$ ,  $-O-Alk-C(NH_2):NOH$ ,  $-O-Alk-NR_5R_6$ ,  $-O-Alk-CN$ ,  $-O-(CH_2)_n-Ph$ ,  $-O-Alk-CO-NR_5R_6$ ,  $-CO-NH-(CH_2)_m-COOR_7$ ,  $-CO-NH-Alk$
- $R_2$  represents H,  $(C_1-C_5)$ alkyl,  $(C_1-C_5)$ alkyl halide,  $(C_3-C_6)$ cycloalkyl or phenyl which is optionally substituted,
- $A$  represents  $-CO-$ ,  $-SO-$  or  $-SO_2-$ ,

- $R_3$  and  $R_4$  which are identical or different, each represent H,  $(C_1-C_5)$ alkoxy, amino, carboxyl,  $(C_2-C_6)$ -alkoxycarbonyl, -OH, nitro, hydroxyamino, -Alk-COOR<sub>7</sub>, -NR<sub>5</sub>R<sub>6</sub>, -NH-Alk-COOR<sub>7</sub>, -NH-COO-Alk, -N(R<sub>11</sub>)-SO<sub>2</sub>-Alk-NR<sub>9</sub>R<sub>10</sub>, -N(R<sub>11</sub>)-SO<sub>2</sub>-Alk, -N(R<sub>11</sub>)-Alk-NR<sub>5</sub>R<sub>6</sub>, -N(R<sub>11</sub>)-CO-Alk-NR<sub>9</sub>R<sub>10</sub>, -N(R<sub>11</sub>)-CO-Alk, -N(R<sub>11</sub>)-CO-CF<sub>3</sub>, -NH-Alk-HetN, -O-Alk-NR<sub>9</sub>R<sub>10</sub>, -O-Alk-CO-NR<sub>5</sub>R<sub>6</sub>, -O-Alk-HetN, or  $R_3$  and  $R_4$  form together a 5- to 6-membered unsaturated heterocycle,
10. optionally in the form of one of their pharmaceutically acceptable salts.